

Applicants : Virginia M. Litwin, et al.
U.S. Serial No.: 09/891,062
Filed : June 25, 2001
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Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of the claims in the application.

In the Claims

Please amend as follows:

Claims 1-39 (Cancelled).

a!
contd.

--40. (Amended) A method of inhibiting HIV-1 infection of a CD4+ cell which comprises contacting the CD4+ cell with an amount of a monoclonal antibody or portion thereof effective to (a) specifically inhibit 67% or greater of fusion of a CD4+ PM-1 cell to a HeLa cell expressing envelope glycoprotein from HIV-1_{JR-FL}, and (b) inhibit 18% or less of fusion of a CD4+ SUP-T1 cell to a HeLa cell expressing envelope protein from HIV-1_{LAI}, wherein the antibody (i) does not crossreact with HIV-1 envelope glycoprotein or CD4; (ii) reacts with an antigen on the surface of a PM-1 cell, and (iii) does not react with an antigen on the surface of a SUP-T1 cell, and (iv) is at least as active as monoclonal antibody PA-7 in inhibiting fusion as recited in (a) above and less active than monoclonal antibody PA-6 in inhibiting fusion as recited in (b) above, so as to thereby inhibit HIV-1 infection of the CD4+ cell.--

--41. (Previously presented) The method of claim 40, wherein the monoclonal antibody is a chimeric monoclonal antibody.--

--42. (Previously presented) The method of claim 40, wherein the monoclonal antibody is humanized.--

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--43. (Previously presented) The method of claim 40, wherein the monoclonal antibody is a human monoclonal antibody.--

--44. (Previously presented) The method of claim 40, wherein the portion of the monoclonal antibody is a single chain antibody or an antigen binding fragment.--

--45. (Previously presented) The method of claim 40, wherein the monoclonal antibody is labeled with a detectable marker.--

al
--46. (Previously presented) The method of claim 45, wherein the detectable marker is a radioactive isotope, enzyme, dye or biotin.--

cond.
--47. (Previously presented) The method of claim 40, wherein the CD4+ cell is present in the subject and the contacting is effected by administering the monoclonal antibody or portion thereof to the subject.--
